



INTERNATIONAL
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CERTIFICATE OF ACCREDITATION

This is to attest that

UL INTERNATIONAL POLSKA SP. Z O.O.

RÓWNOLEGŁA 4
WARSZAWA 02-235, POLAND

Testing Laboratory TL-863

has met the requirements of AC89, *IAS Accreditation Criteria for Testing Laboratories* and has demonstrated compliance with ISO/IEC Standard 17025:2017, *General requirements for the competence of testing and calibration laboratories*. This organization is accredited to provide the services specified in the scope of accreditation.

Effective Date July 22, 2025



International Accreditation Service
Issued under the authority of IAS management

Visit www.iasonline.org for current accreditation information.

SCOPE OF ACCREDITATION

International Accreditation Service, Inc.

3060 Saturn Street, Suite 101, Brea, California 92821, U.S.A. | www.iasonline.org

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www.ul.com

Contact Name Marcin Wagner

Contact Phone +48 22 336 3367

Accredited to ISO/IEC 17025:2017

Effective Date July 22, 2025

| Electrical | |
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| ANSI/AAMI ES60601-1 IEC/EN 60601-1 | Medical electrical equipment – Part 1: General requirements for basic safety and essential performance |
| ANSI AAMI HA60601-1-11 | Medical Electrical Equipment -- Part 1-11: General requirements for basic safety and essential performance -- Collateral Standard: Requirements for medical electrical equipment and medical electrical equipment and medical electrical systems used in the home healthcare environment |
| ANSI AAMI IEC 60601-1-12 | Medical electrical equipment - Part 1-12: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems intended for use in the emergency |
| ANSI AAMI IEC 60601-2-25 | Medical electrical equipment - Part 2-25: Particular requirements for the basic safety and essential performance of electrocardiographs |
| ANSI AAMI IEC 60601-2-27 | Medical electrical equipment - Part 2-27: Particular requirements for the basic safety and essential performance of electrocardiographic monitoring equipment |
| ANSI AAMI IEC 60601-2-47 | Medical electrical equipment - Part 2-47: Particular requirements for the basic safety and essential performance of ambulatory electrocardiographic systems |
| ANSI UL 61010-1 | Standard for Safety for Electrical Equipment For Measurement, Control and Laboratory Use; Part 1: General Requirements |
| ANSI/UL/IEC/EN 61010-2-201 | Safety requirements for electrical equipment for measurement, control, and laboratory use – Part 2-201: Particular requirements for control equipment |
| IEC 61010-1 Edition 3.1 2017-01 | Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements |
| IEC/EN 60601-1-6 | Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability (applicable test standard IEC 60601-1) |
| IEC/EN 60601-1-11 | Medical electrical equipment - Part 1-11: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems used in the home healthcare environment |

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| IEC/EN 60601-1-12 | Medical electrical equipment - Part 1-12: General requirements for basic safety and essential performance - Collateral Standard: Requirements for medical electrical equipment and medical electrical systems intended for use in the emergency medical services environment |
| IEC/EN 60601-2-24 | Medical electrical equipment - Part 2-24: Particular requirements for the basic safety and essential performance of infusion pumps and controllers |
| IEC/EN 60601-2-25 | Medical electrical equipment - Part 2-25: Particular requirements for the basic safety and essential performance of electrocardiographs |
| IEC/EN 60601-2-27 | Medical electrical equipment - Part 2-27: Particular requirements for the basic safety and essential performance of electrocardiographic monitoring equipment |
| IEC/EN 60601-2-47 | Medical electrical equipment - Part 2-47: Particular requirements for the basic safety and essential performance of ambulatory electrocardiographic systems |
| IEC/EN 60601-2-49 | Medical electrical equipment - Part 2-49: Particular requirements for the basic safety and essential performance of multifunction patient monitors |
| IEC/EN 60601-2-52 | Medical electrical equipment – Part 2-52: Particular requirements for the basic safety and essential performance of medical beds |
| IEC/EN 60950-1 | Information technology equipment – Safety – Part 1: General requirements |
| IEC/EN 62368-1 | Audio/video, information and communication technology equipment – Part 1: Safety requirements |
| IEC/EN 80601-2-49 | Medical electrical equipment - Part 2-49: Particular requirements for the basic safety and essential performance of multifunction patient monitors |
| IEC/EN 80601-2-60 | Medical electrical equipment – Part 2-60: Particular requirements for the basic safety and essential performance of dental equipment |
| IEC/EN 80601-2-61 | Medical electrical equipment - Part 2-61: Particular requirements for basic safety and essential performance of pulse oximeter equipment |
| UL/IEC/EN 61010-1 | Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements |
| UL/IEC/EN 61010-2-010 | Safety requirements for electrical equipment for measurement, control and laboratory use – Part 2-010: Particular requirements for laboratory equipment for the heating of materials |
| UL/IEC/EN 61010-2-030 | Safety requirements for electrical equipment for measurement, control, and laboratory use – Part 2-030: Particular requirements for equipment having testing or measuring circuits |
| IEC 61010-2-081 | Safety requirements for electrical equipment for measurement, control and laboratory use - Part 2-081: Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes |

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| IEC 61010-2-101 | Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-101: Particular requirements for in vitro diagnostic (IVD) medical equipment |
| Battery | |
| 16 CFR 1263 | Button cell and coin batteries and consumer products containing such batteries |
| IEC/EN 60086-4 | Primary batteries – Part 4: Safety of lithium batteries |
| IEC/EN 62133 | Secondary cells and batteries containing alkaline or other non-acid electrolytes – Safety requirements for portable sealed secondary cells, and for batteries made from them, for use in portable applications |
| IEC/EN 62133-1 | Secondary cells and batteries containing alkaline or other non-acid electrolytes - Safety requirements for portable sealed secondary cells, and for batteries made from them, for use in portable applications - Part 1: Nickel systems |
| IEC/EN 62133-2 | Secondary cells and batteries containing alkaline or other non-acid electrolytes – Safety requirements for portable sealed secondary cells, and for batteries made from them, for use in portable applications – Part 2: Lithium systems |
| EN 50604-1 | Secondary lithium batteries for light EV (electric vehicle) applications - Part 1: General safety requirements and test methods |
| IEC/EN 62281 | Safety of primary and secondary lithium cells and batteries during transport |
| IEC 62660-3 | Secondary lithium-ion cells for the propulsion of electric road vehicles - Part 3: Safety requirements |
| IEC 62619 | Secondary cells and batteries containing alkaline or other non-acid electrolytes - Safety requirements for secondary lithium cells and batteries, for use in industrial applications Exclusion: Clause 7.2.3.3 |
| IEC 62841-1 | Electric motor-operated hand-held tools, transportable tools and lawn and garden machinery - Safety - Part 1: General requirements Only: Annex K and Annex L |
| IEC 60335-1 | Household and similar electrical appliances - Safety - Part 1: General requirements Only: Clause 21.1 and Annex B |
| IEC 63370 | Lithium-ion batteries and charging systems - Safety |
| UL 2271 | Batteries for Use In Light Electric Vehicle (LEV) Applications |
| UL 2272 | Electrical Systems for Personal E- Mobility Devices |
| UL 2849 | Electrical Systems for eBikes |
| UL 4200A | Products incorporating button batteries or coin cell batteries |

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| UN ST/SG/AC.10/11 | Recommendations on the Transport of Dangerous Goods, Manual of Tests and Criteria. Clause 38.3 Lithium metal and lithium ion batteries |
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